

IN THE CLAIMS

Claim 1 (currently amended) A stent delivery system comprising:

a catheter having a guidewire lumen at least partially therethrough, and having a fluid pressurizing lumen therein separate from said guidewire lumen, said fluid pressurizing lumen extending between a proximal end thereof and a fluid opening at a distal end of said catheter;

said catheter having a compressed stent retention section which extends at least a compressed stent length in a first direction along said catheter from a stent ~~exp~~ plunger engaged with said catheter at a location near a distal end of said catheter, said catheter having a sheath retraction section which extends from said stent ~~exp~~ plunger in a second direction which is opposite said first direction to a fixed seal mount fixed to said catheter, said catheter further containing a fluid receiving chamber section containing said fluid opening, said fluid receiving chamber section extends from said fixed seal mount in said second direction to a maximum fluid receiving chamber extension length;

a stent containment sheath having a movable seal mount near an end thereof, wherein said stent containment sheath in a pre stent deployment position is positioned surrounding a portion of a distal end of said catheter including said compressed stent retention section, said stent ~~exp~~ plunger, said sheath retraction section, said fixed seal mount, and a minimum fluid receiving chamber extension length, wherein said stent containment sheath in a post stent deployment position is positioned surrounding a portion of a distal end of said catheter including said stent ~~exp~~ plunger, said sheath retraction section, said fixed seal mount, and a substantial portion of said maximum fluid receiving chamber extension length, wherein a stent retention portion of said stent containment sheath is made of a first material having a lubricious inner surface suitable for easy release of a stent contained therein, wherein a stent retraction portion of said stent containment sheath is made of a second material different from said first material having a smooth inner surface against which a first flexible seal structure of a fluid receiving chamber seals;

wherein said first flexible seal structure is disposed engaged with said fixed seal mount to flexibly seal between said catheter and said stent containment sheath

at a first end of said fluid receiving chamber, said first flexible seal structure maintains engagement with said fixed seal mount as said fluid receiving chamber is pressurized and said stent containment sheath moves with respect to said catheter; and

a second flexible seal structure disposed engaged with said movable seal mount and flexibly sealing between said catheter and said stent containment sheath at an end opposite said first end of a fluid receiving section and maintaining engagement with said movable seal mount as said fluid receiving chamber is pressurized and said stent containment sheath moves with respect to said catheter.

1 Claim 2 (original) A stent delivery system as in Claim 1, further comprising:
2 an anti-kinking spacer loosely contained within said stent containment
3 sheath and outside said sheath retraction section of said catheter and extending
4 substantially between ends of said sheath retraction section and sized to substantially
5 interfere with the kinking of said stent containment sheath at a location adjacent to said
6 anti-kinking spacer when said stent containment sheath containing a portion of said
7 catheter is bent.

1 Claim 3 (original) A stent delivery system as in Claim 2,
2 wherein said anti-kinking spacer is a helical spring.

1 Claim 4 (original) A stent delivery system as in Claim 3,
2 wherein said anti-kinking spacer is a helical spring having a substantially planar
3 coil shape.

1 Claim 5 (original) A stent delivery system as in Claim 4,
2 wherein said anti-kinking spacer is a helical spring having a substantially planar
3 coil shape whose thickness larger near the central longitudinal axis of the helix and tapers
4 to a smaller thickness near its outer edge.

1 Claim 6 (currently amended) A stent delivery system as in Claim 2,
2 wherein said anti-kinking spacer ~~is a~~ is a series of stacked rings.

1 Claim 7 (original) A stent delivery system as in Claims 1, 2, 3, 4, 5, or 6,
2 wherein ~~and~~ an inside diameter of said stent containment sheath opposite said
3 compressed stent retention section and the inside diameter of said stent containment
4 sheath opposite said retraction section and said fluid receiving chamber are the
5 substantially the same.

1 Claim 8 (original) A stent delivery system as in Claim 7,
2 wherein said catheter has fixed to it a backstop which prevents fluid from being
3 released from the fluid receiving chamber by the stent containment sheath moving so far
4 with respect to the catheter that said fluid receiving chamber is no longer sealed by said
5 first flexible seal structure.

1 Claim 9 (original) A stent delivery system as in Claim 1, 2, 3, 4, 5, or 6,
2 wherein ~~and~~ an inside diameter of said stent containment sheath opposite said
3 compressed stent retention section and the inside diameter of said stent containment
4 sheath opposite said retraction section and said fluid receiving chamber are substantially
5 different.

1 Claim 10 (original) A stent delivery system as in Claim 9,
2 wherein said catheter has fixed to it a backstop which prevents fluid from being
3 released from the fluid receiving chamber by the stent containment sheath moving so far
4 with respect to the catheter that said fluid receiving chamber is no longer sealed by said
5 first flexible seal structure.

1 Claim 11 (currently amended) A stent graft delivery system comprising:
2 a catheter having a guidewire lumen at least partially therethrough, and
3 having a fluid pressurizing lumen therein separate from said guidewire lumen, said fluid
4 pressurizing lumen extending between a proximal end thereof and a fluid opening at a
5 distal end of said catheter;

6 said catheter having a compressed stent graft retention section which
7 extends at least a compressed stent graft length in a first direction along said catheter
8 from a stent graft ~~exp~~ plunger engaged with said catheter at a location near a distal end of
9 said catheter, said catheter having a sheath retraction section which extends from said
10 stent graft ~~exp~~ plunger in a second direction which is opposite said first direction to a
11 fixed seal mount fixed to said catheter, said catheter further containing a fluid receiving
12 chamber section containing said fluid opening, said fluid receiving chamber section
13 extends from said fixed seal mount in said second direction to a maximum fluid receiving
14 chamber extension length;

15 a stent graft containment sheath having a movable seal mount near an end
16 thereof, wherein said stent graft containment sheath in a pre stent graft deployment
17 position is positioned surrounding a portion of a distal end of said catheter including said
18 compressed stent graft retention section, said stent graft ~~exp~~ plunger, said sheath
19 retraction section, said fixed seal mount, and a minimum fluid receiving chamber
20 extension length, wherein said stent graft containment sheath in a post stent graft
21 deployment position is positioned surrounding a portion of a distal end of said catheter
22 including said stent graft ~~exp~~ plunger, said sheath retraction section, said fixed seal
23 mount, and a substantial portion of said maximum fluid receiving chamber extension
24 length, wherein a stent graft retention portion of said stent graft containment sheath is
25 made of a first material having a lubricious inner surface suitable for easy release of a
26 stent graft contained therein, wherein a stent graft retraction portion of said stent graft
27 containment sheath is made of a second material different from said first material having
28 a smooth inner surface against which a first flexible seal structure of a fluid receiving
29 chamber seals;

30 wherein said first flexible seal structure is disposed engaged with said
31 fixed seal mount to flexibly seal between said catheter and said stent graft containment
32 sheath at a first end of said fluid receiving chamber, said first flexible seal structure
33 maintains engagement with said fixed seal mount as said fluid receiving chamber is
34 pressurized and said stent graft containment sheath moves with respect to said catheter;
35 and

36 a second flexible seal structure disposed engaged with said movable seal
37 mount and flexibly sealing between said catheter and said stent graft containment sheath
38 at an end opposite said first end of a fluid receiving section and maintaining engagement
39 with said movable seal mount as said fluid receiving chamber is pressurized and said
40 stent graft containment sheath moves with respect to said catheter.

1 Claim 12 (original) A stent graft delivery system as in Claim 11, further comprising:
2 an anti-kinking spacer loosely contained within said stent graft
3 containment sheath and outside said sheath retraction section of said catheter and
4 extending substantially between ends of said sheath retraction section and sized to
5 substantially interfere with the kinking of said stent graft containment sheath at a location
6 adjacent to said anti-kinking spacer when said stent graft containment sheath containing a
7 portion of said catheter is bent.

1 Claim 13 (original) A stent graft delivery system as in Claim 12,
2 wherein said anti-kinking spacer is a helical spring.

1 Claim 14 (original) A stent graft delivery system as in Claim 13,
2 wherein said anti-kinking spacer is a helical spring having a substantially planar
3 coil shape.

1 Claim 15 (original) A stent graft delivery system as in Claim 14,
2 wherein said anti-kinking spacer is a helical spring having a substantially planar
3 coil shape whose thickness larger near the central longitudinal axis of the helix and tapers
4 to a smaller thickness near its outer edge.

1 Claim 16 (currently amended) A stent graft delivery system as in Claim 12,
2 wherein said anti-kinking spacer ~~is a~~ is a series of stacked rings.

1 Claim 17 (original) A stent graft delivery system as in Claims 11, 12, 13, 14, 15, or 16,

2 wherein ~~and~~ an inside diameter of said stent graft containment sheath opposite
3 said compressed stent graft retention section and the inside diameter of said stent graft
4 containment sheath opposite said retraction section and said fluid receiving chamber are
5 the substantially the same.

1 Claim 18 (original) A stent graft delivery system as in Claim 17,
2 wherein said catheter has fixed to it a backstop which prevents fluid from being
3 released from the fluid receiving chamber by the stent graft containment sheath moving
4 so far with respect to the catheter that said fluid receiving chamber is no longer sealed by
5 said first flexible seal structure.

1 Claim 19 (original) A stent graft delivery system as in Claim 11, 12, 13, 14, 15, or 16,
2 wherein ~~and~~ an inside diameter of said stent graft containment sheath opposite
3 said compressed stent graft retention section and the inside diameter of said stent graft
4 containment sheath opposite said retraction section and said fluid receiving chamber are
5 substantially different.

1 Claim 20 (original) A stent graft delivery system as in Claim 19,
2 wherein said catheter has fixed to it a backstop which prevents fluid from being
3 released from the fluid receiving chamber by the stent graft containment sheath moving
4 so far with respect to the catheter that said fluid receiving chamber is no longer sealed by
5 said first flexible seal structure.

1 Claim 21 (currently amended) A stent delivery system comprising:
2 a catheter having a guidewire lumen at least partially therethrough, and
3 having a fluid pressurizing lumen therein separate from said guidewire lumen, said fluid
4 pressurizing lumen extending between a proximal end thereof and a fluid opening at a
5 distal end of said catheter;
6 said catheter having a compressed stent retention section which extends at
7 least a compressed stent length in a first direction along said catheter from a stent ~~exp~~
8 plunger engaged with said catheter at a location near a distal end of said catheter, said

9 catheter having a sheath retraction section which extends from said stent ~~exp~~ plunger in a
10 second direction which is opposite said first direction to a fixed seal mount fixed to said
11 catheter, said catheter further containing a fluid receiving chamber section containing
12 said fluid opening, said fluid receiving chamber section extends from said fixed seal
13 mount in said second direction to a maximum fluid receiving chamber extension length;
14 a stent containment sheath having a movable seal mount near an end
15 thereof, wherein said stent containment sheath in a pre stent deployment position is
16 positioned surrounding a portion of a distal end of said catheter including said
17 compressed stent retention section, said stent ~~exp~~ plunger, said sheath retraction section,
18 said fixed seal mount, and a minimum fluid receiving chamber extension length, wherein
19 said stent containment sheath in a post stent deployment position is positioned
20 surrounding a portion of a distal end of said catheter including said stent ~~exp~~ plunger,
21 said sheath retraction section, said fixed seal mount, and a substantial portion of said
22 maximum fluid receiving chamber extension length;
23 a first flexible seal structure disposed engaged with said fixed seal mount
24 and flexibly sealing between said catheter and said stent containment sheath at a first end
25 of said fluid receiving chamber, said first flexible seal structure maintains engagement
26 with said fixed seal mount as said fluid receiving chamber is pressurized and said stent
27 containment sheath moves with respect to said catheter;
28 a second flexible seal structure disposed engaged with said movable seal
29 mount and flexibly sealing between said catheter and said stent containment sheath at an
30 end opposite said first end of a fluid receiving section and maintaining engagement with
31 said movable seal mount as said fluid receiving chamber is pressurized and said stent
32 containment sheath moves with respect to said catheter; and.
33 an anti-kinking spacer loosely contained within said stent containment
34 sheath and outside said sheath retraction section of said catheter and extending
35 substantially between ends of said sheath retraction section and sized to substantially
36 interfere with the kinking of said stent containment sheath at a location adjacent to said
37 anti-kinking spacer when said stent containment sheath containing a portion of said
38 catheter is bent.

1 Claim 22 (currently amended) A stent graft delivery system comprising:
2 a catheter having a guidewire lumen at least partially therethrough, and
3 having a fluid pressurizing lumen therein separate from said guidewire lumen, said fluid
4 pressurizing lumen extending between a proximal end thereof and a fluid opening at a
5 distal end of said catheter;
6 said catheter having a compressed stent graft retention section which
7 extends at least a compressed stent graft length in a first direction along said catheter
8 from a stent graft ~~exp~~ plunger engaged with said catheter at a location near a distal end of
9 said catheter, said catheter having a sheath retraction section which extends from said
10 stent graft ~~exp~~ plunger in a second direction which is opposite said first direction to a
11 fixed seal mount fixed to said catheter, said catheter further containing a fluid receiving
12 chamber section containing said fluid opening, said fluid receiving chamber section
13 extends from said fixed seal mount in said second direction to a maximum fluid receiving
14 chamber extension length;
15 a stent graft containment sheath having a movable seal mount near an end
16 thereof, wherein said stent graft containment sheath in a pre stent graft deployment
17 position is positioned surrounding a portion of a distal end of said catheter including said
18 compressed stent graft retention section, said stent graft ~~exp~~ plunger, said sheath
19 retraction section, said fixed seal mount, and a minimum fluid receiving chamber
20 extension length, wherein said stent graft containment sheath in a post stent graft
21 deployment position is positioned surrounding a portion of a distal end of said catheter
22 including said stent graft ~~exp~~ plunger, said sheath retraction section, said fixed seal
23 mount, and a substantial portion of said maximum fluid receiving chamber extension
24 length;
25 a first flexible seal structure disposed engaged with said fixed seal mount
26 and flexibly sealing between said catheter and said stent graft containment sheath at a
27 first end of said fluid receiving chamber, said first flexible seal structure maintains
28 engagement with said fixed seal mount as said fluid receiving chamber is pressurized and
29 said stent graft containment sheath moves with respect to said catheter;
30 a second flexible seal structure disposed engaged with said movable seal
31 mount and flexibly sealing between said catheter and said stent graft containment sheath

32 at an end opposite said first end of a fluid receiving section and maintaining engagement
33 with said movable seal mount as said fluid receiving chamber is pressurized and said
34 stent graft containment sheath moves with respect to said catheter; and.

35 an anti-kinking spacer loosely contained within said stent graft
36 containment sheath and outside said sheath retraction section of said catheter and
37 extending substantially between ends of said sheath retraction section and sized to
38 substantially interfere with the kinking of said stent graft containment sheath at a location
39 adjacent to said anti-kinking spacer when said stent graft containment sheath containing a
40 portion of said catheter is bent.

1 Claim 23 (original) The stent delivery system as in Claim 1 or 11
2 wherein a wall thickness of said first material of said stent retention portion is
3 different than a wall thickness of said second material of said stent retraction portion.

1 Claim 24 (currently amended) The system according to Claim 11 wherein said
2 stent graft assembly is a self-expanding stent graft-assembly.

1 Claim 25 (currently amended) A method for hydraulically retracting a stent
2 containment sheath comprising the steps of:
3 providing a catheter having fixed seal fixed to a fixed seal mount thereon, with a
4 fluid receiving chamber section on one side of said fixed seal and an anti kinking spacer
5 on a second side of said with a plunger ~~cup~~ disposed at the end of said antikinking spacer
6 opposite the fixed seal with a stent in a compressed pre deployment position disposed
7 around ~~the stent~~ at a stent retention section of the catheter beyond the plunger ~~cup~~;
8 surrounding a portion of a distal end of said catheter with a containment sheath
9 such sheath containing said fixed seal and said fixed seal mount and said antikinking
10 spacer and said plunger ~~cup~~ and said stent in said pre deployment position, said
11 containment sheath being sized to seal against said fixed seal of said catheter and
12 including a movable seal which moves with the containment catheter and seals against
13 said catheter to establish a fluid receiving chamber between the catheter, the containment
14 sheath and the fixed seal and the movable seal; and

15 injecting fluid into a lumen of said catheter in communication with a fluid
16 opening in said fluid receiving chamber, such pressurization causing said retraction
17 sheath to retract with respect to said catheter and uncover the stent for deployment.

1 Claim 26 (previously amended) The method of Claims 24,
2 wherein the stent containment sheath is constructed from at least two different
3 diameter tubes.

1 Claim 27 (currently amended) The method of Claim 24,
2 wherein the stent containment sheath is constructed from at least two different
3 ~~materials~~ material having substantially different surface lubricity.

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